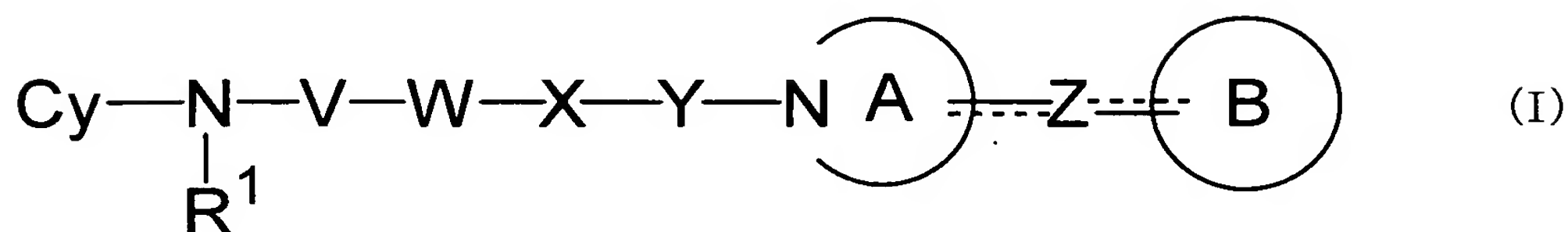


CLAIMS

1. A compound represented by Formula (1):

[Formula 1]



wherein Cy is an aromatic hydrocarbon group which may be substituted, or an aromatic heterocyclic group which may be substituted; R¹ is a hydrogen atom or a hydrocarbon group which may be substituted; V is -C(O)-, -S(O)-, or -S(O)₂-; W is -N(R²)-, -O-, or a bond (wherein R² is a hydrogen atom or a hydrocarbon group which may be substituted); X is alkylene which may be substituted; Y is -C(O)-, -S(O)-, or -S(O)₂-; Z is a bond, a chain hydrocarbon group which may be substituted, or -N=; ring A is a non-aromatic nitrogen-containing heterocyclic ring which may be substituted; and ring B is a nitrogen-containing heterocyclic group which may be substituted;

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is each independently a single bond or a double bond; R¹ and R² may be bonded to each other to form a non-aromatic nitrogen-containing heterocyclic ring which may be substituted; and R² may be bonded to a substituent of X to form a non-aromatic nitrogen-containing heterocyclic ring which may be substituted,

or a salt thereof.

2. A prodrug of the compound according to claim 1.

3. The compound according to claim 1, wherein Cy is
5 phenyl which may be substituted, or a 5- to 6-membered
aromatic monocyclic heterocyclic group which may be
substituted.

4. The compound according to claim 1, wherein Cy is
10 phenyl which may be substituted with a halogen atom.

5. The compound according to claim 1, wherein R^1 is a
hydrogen atom.

15 6. The compound according to claim 1, wherein V is -
C(O)-.

7. The compound according to claim 1, wherein W is -
N(R^2)-.

20

8. The compound according to claim 1, wherein X is C_{1-4}
alkylene which may be substituted with a hydrocarbon group
which may be substituted, an aromatic heterocyclic group
which may be substituted, a hydroxyl group which may be
25 substituted, amino which may be substituted, carbamoyl
which may be substituted or carboxyl which may be

esterified.

9. The compound according to claim 1, wherein X is methylene which may be substituted with a hydrocarbon group
5 which may be substituted or an aromatic heterocyclic group which may be substituted.

10. The compound according to claim 1, wherein Y is -
C(O)-.
10

11. The compound according to claim 1, wherein -W-X-Y-
is an amino acid residue.

12. The compound according to claim 1, wherein ring A is
15 a piperidine ring which may be substituted, or a piperazine ring which may be substituted.

13. The compound according to claim 1, wherein ring B is
a monocyclic nitrogen-containing heterocyclic ring which
20 may be substituted.

14. The compound according to claim 13, wherein the
monocyclic nitrogen-containing heterocyclic ring is a
piperidine ring, a piperazine ring, a morpholine ring, an
25 imidazoline ring, a pyrrolidine ring, a pyridine ring, an
imidazole ring, or a thiazoline ring.

15. The compound according to claim 1, wherein ring B is a fused nitrogen-containing heterocyclic ring which may be substituted.

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16. The compound according to claim 15, wherein the fused nitrogen-containing heterocyclic ring is a fused pyridine ring, a fused imidazole ring, a fused pyrazole ring, or a fused thiazoline ring.

10

17. The compound according to claim 1, wherein Z is a bond or C₁₋₆ alkylene.

18. A compound selected from the group consisting of N-(4-chlorophenyl)-N'-((1R)-2,2-dimethyl-1-((4-(5-methyl-3-oxo-1H-imidazo[1,5-c]imidazol-2(3H)-yl)-1-piperazinyl)carbonyl)propyl)urea, N-(4-chlorophenyl)-N'-(2-ethyl-2-hydroxy-1-((4-(5-methyl-3-oxo-1H-imidazo[1,5-c]imidazol-2(3H)-yl)-1-piperazinyl)carbonyl)butyl)urea, N-(4-chlorophenyl)-N'-((1S)-2-methyl-1-((4-(5-methyl-3-oxo-1H-imidazo[1,5-c]imidazol-2(3H)-yl)-1-piperazinyl)carbonyl)-2-(methylthio)propyl)urea, and N-(4-chlorophenyl)-N'-(2-methoxy-2-methyl-1-((4-(5-methyl-3-oxo-1H-imidazo[1,5-c]imidazol-2(3H)-yl)-1-piperazinyl)carbonyl)propyl)urea, or a salt thereof.

19. A pharmaceutical composition comprising the compound according to claim 1 or 2.

20. The pharmaceutical composition according to claim 19,
5 which is an anticoagulant.

21. The pharmaceutical composition according to claim 19,
which is an activated blood coagulation factor X inhibitor.

10 22. The pharmaceutical composition according to claim 19,
which is a prophylactic and/or therapeutic agent for
myocardial infarction, cerebral infarction, deep vein
thrombosis, pulmonary thromboembolism, or arteriosclerosis
obliterans.

15

23. The pharmaceutical composition according to claim 19,
which is a prophylactic and/or therapeutic agent for
economy-class syndrome, thromboembolism during and post
operation, or the secondary onset of deep vein thrombosis.

20

24. A method of inhibiting blood coagulation in mammal
which comprises administering an effective amount of the
compound according to claim 1 or a prodrug thereof to the
mammal.

25

25. A method of inhibiting activated blood coagulation

factor X in mammal which comprises administering an effective amount of the compound according to claim 1 or a prodrug thereof to the mammal.

5 26. A method of preventing and/or treating myocardial infarction, cerebral infarction, deep vein thrombosis, pulmonary thromboembolism or arteriosclerosis obliterans in mammal which comprises administering an effective amount of the compound according to claim 1 or a prodrug thereof to
10 the mammal.

15 27. Use of the compound according to claim 1 or a prodrug thereof, for the manufacture of a medicine for inhibiting blood coagulation.

20 28. Use of the compound according to claim 1 or a prodrug thereof, for the manufacture of a medicine for inhibiting activated blood coagulation factor X.

25 29. Use of the compound according to claim 1 or a prodrug thereof, for the manufacture of a medicine for preventing and/or treating myocardial infarction, cerebral infarction, deep vein thrombosis, pulmonary thromboembolism, or arteriosclerosis obliterans.